Research Bacteriology -Bldg. 54 - Rm. 16



VETERANS ADMINISTRATION

HOSPITAL
Hines, Illinois

YOUR FILE REFERENCE:

September 19, 1952

IN REPLY REFER TO:

Dr. Joshua Lederberg Department of Genetics College of Agriculture Madison 6, Wisconsin

Dear Dr. Lederberg:

We have used no very precise methods in preparing the McLeod's synthetic medium except for using reagents of highest purity for our work with M N Q. The composition of the medium was as follows:

Ammonium sulfate	4.72 grams
Asparagine	2.0 "
Sodium chloride	5.0 "
Glucose	2.0 "
$\mathbb{KH}_2\mathbb{P}_4 \left(\underline{\mathbf{M}}\right)$	
5	100. ml.
Va OH (N)	16.4 ml. (or to pH 7.8)
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0.5% solutions of MgCl2, Fe Cl2 and Ca Cl2 were freshly prepared separately, then mixed, and 1.0 ml of the mixture was added to the medium.

Distilled water

1000 ml.

Chemicals were weighed only roughly on a triple beam balance accurate to 0.01 gram, even for the $\rm M/5~KH_2~PO_4$ solution.

pH should be about 7.8 before autoclaving, using the amount of NaOH designated; if not, NaOH was added to this point.

pH after autoclaving at 12 lbs. for 20 minutes, 7.6.

Medium was distributed by hand-pipetting with 25 ml measuring pipettes in 24.5 ml. amounts into 8^n x 1^n test tubes.

Difco nutrient agar, (B 69), was used for streaking cultures and for slants.

I trust that your efforts will meet with more success this time.

Sincerely colivel